

Start Time	Discussion
8:00 a.m.	Sarah Acaster, Andrew Lloyd, Oxford Outcomes Ltd, Patient Reported Outcomes
8:20 a.m.	Q&A
8:30 a.m.	Zeeshan Butt, Bryce Reeve, Northwestern University , Department of Medical Social Sciences/ University of North Carolina-Chapel Hill , Department of Health Policy & Management
8:50 a.m.	Q&A
9:00 a.m.	Group Discussion —Research Teams, External Invitees, & Workgroup Members
10:00 a.m.	Break
10:15 a.m.	Discussion of Report Content —Research Teams, External Invitees, & Workgroup Members
12:00 p.m.	Ethan Basch, Mary Tinetti, Closing Remarks
12:15 p.m.	Lunch (<i>Ravenhurst Room</i>)
1:00 p.m.	Adjourn

- Standards or principles?
- “Minimum”
- Levels: supra- vs. subordinate
- Prioritization
- Actionable
- Strength of evidence
- How to communicate?
- Knowledge gaps

**Oxford Outcomes
Northwestern University/UNC Chapel Hill**

- Is PROM use for PCOR different in any way from PROM use in research generally?
 - When would this be so?
 - How do the proposed standards for PRO use in PCOR differ from other current standards for PRO use? Should PCOR-specific standards be adopted more generally?
- Scope for this work was limited to PROM use in research settings. How would the standards proposed here relate to PROM use in clinical contexts, and could they support the PCORI research priority on improving healthcare systems?
- What are methodological and logistical challenges of applying the proposed standards in "real-world" or non-experimental settings?
- What are examples of PROMs that adhere to the proposed standards?